1. (Previously presented) A compound according to the structure:

where X is (CH<sub>2</sub>)<sub>m</sub>COR

R is a  $C_1$  to  $C_5$  alkyl group optionally substituted with at least one halogen group; and m is from 0-5, and pharmaceutically acceptable salts, solvates or polymorphs thereof.

- 2. (Original) The compound according to claim 1 wherein F is a  $C_1$  to  $C_5$  alkyl group optionally substituted with at least one halogen group and m is from 0-:...
- 3. (Original) The compound according to claim 1 wherein R is methyl, ethyl, propyl, iso-propyl, butyl, iso-butyl, neo-pentyl or CH<sub>2</sub>CH<sub>2</sub>F; and m is 0.
- 4. (Original) The compound according to claim 3 wherein n is 0 and R is methyl, ethyl or  $CH_2CH_2F$ .
- 5. (Original) The compound according to claim 4 wherein R is methyl.
- 6. (Original) The compound according to claim 4 wherein R is ethyl.
- 7. (Original) The compound according to claim 4 wherein R is CH<sub>2</sub>CH<sub>2</sub>F.

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- 8. (Original) The compound according to claim 1 wherein F is CH<sub>2</sub>CHF<sub>2</sub>.
- 9. (Original) The compound according to claim 1 wherein I is CH<sub>2</sub>CF<sub>3</sub>.
- 10. (Original) A compound according to claim 1, wherein F is a C<sub>1</sub> to C<sub>5</sub> alkyl, which may be unsubstituted or substituted with at least one F group.
- 11. (Previously presented) The compound according to claim 9 wherein R is methyl, ethyl or CH<sub>2</sub>CH<sub>2</sub>F.
- 12. (Currently amended) A pharmaceutical composition consisting essentially of an effective amount of a compound for alleviating the symptomology of menopause in a patient, said compound having the structure:

R is H-or a C<sub>1</sub> to C<sub>5</sub> alkyl group optionally substituted with at least one halogen group, and m is from 0-5, and pharmaceutically acceptable salts, solvates or polymorphs thereof, optionally in combination with a pharmaceutically acceptable carrier, excipient or additive.

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- 13. (Previously presented) The composition according to claim 12 wherein R is a C<sub>1</sub> to C<sub>5</sub> alkyl group optionally substituted with at least one fluorine group, and m is from 0-2.
- 14. (Original) The composition according to claim 12 wherein R is methyl, ethyl, propyl, isopropyl, butyl, iso-butyl, pentyl, neo-pentyl, CH<sub>2</sub>CH<sub>2</sub>F or CH CF<sub>3</sub> and m is 0.
- 15. (Original) The composition according to claim 14 where in m is 0 and R is methyl, ethyl or CH<sub>2</sub>CH<sub>2</sub>F.
- 16. (Original) The composition according to claim 14 where in R is methyl.
- 17. (Original) The composition according to claim 14 wherein R is ethyl.
- 18. (Original) The composition according to claim 14 when an R is CH<sub>2</sub>CH<sub>2</sub>F.
- 19. (Original) The composition according to claim 12 in topical dosage form.
- 20. (Original) The composition according to claim 12 formulated as a vaginal cream, gel, lotion or suppository.
- 21. (Currently amended) A method for alleviating the symptoms of menopause, comprising administering to a patient in need of therapy a pharmaceutical composition comprising an effective amount of a compound according to the structure:

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R is H or a C<sub>1</sub> to C<sub>5</sub> alkyl group, optionally substituted with at least one halogen group; and m is from 0-5, and pharmaceutically acceptable salts, so vates or polymorphs thereof, optionally in combination with a pharmaceutically acceptable carrier, excipient or additive.

- 22. (Original) The method according to claim 21 wherein R is a  $C_1$  to  $C_5$  alkyl group or a  $CH_2CH_2F$  group; and m is from 0-2.
- 23. (Original) The method according to claim 21 wherein said symptom of menopause is selected from the group consisting of bone loss associated with osteoporosis and vaginal dyspareunia.
- 24. (Original) The method according to claim 21 wherein said symptom of menopause is vaginal dyspareunia and said composition is administered to the patient's vaginal membranes.
- 25. (Original) The method according to claim 22 wherein R is methyl.
- 26. (Original) The method according to claim 22 wherein R is ethyl.
- 27. (Original) The method according to claim 22 wherein R:s CH<sub>2</sub>CH<sub>2</sub>F.

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- 28. (Original) The method according to claim 22 wherein R is CH<sub>2</sub>CHF<sub>2</sub>.
- 29. (Original) The method according to claim 22 wherein R is CH<sub>2</sub>CF<sub>3</sub>.
- 30. (Original) The method according to claim 24 wherein said composition is administered as a vaginal cream, gel, lotion or suppository.
- 31. (Original) The method according to claim 21 wherein said composition is administered within the patient's body from an implant.
- 32. (Original) The method according to claim 21 wherein said symptom of menopause is bone loss associated with osteoporosis and said composition is administered within the patient's body from an implant.
- 33. (Previously presented) A compound according to the structure:

R is H; and

m is from 0, 2, 3, 4, or 5, and pharmaceutically acceptable salts, solvates or polymorphs thereof.

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- 34. (Previously presented) The compound according to claum 33 wherein m is 0.
- 35. (Previously presented) The compound according to claim 33 wherein m is 2, 3, 4, or 5.

The following claims are new:

36. (New) A pharmaceutical composition consisting essentially of an effective amount of a compound for alleviating the symptomology of menopause in a patient, said compound having the structure:

R is H; and

m is from 0, 2, 3, 4, or 5, and pharmaceutically acceptable salts, solvates or polymorphs thereof, optionally in combination with a pharmaceutically acceptable carrier, excipient or additive.

- 37. (New) The composition according to claim 36 wherein in is 0.
- 38. (New) The composition according to claim 36 wherein n is 2, 3, 4, or 5.

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